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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/593,034	09/15/2006	Keiko Shimamoto	47233-5007-00 (230642) 5739	
	7590 08/02/201 DDLE & REATH (DC)	EXAMINER		
1500 K STREE		BARKER, MICHAEL P		
SUITE 1100 WASHINGTON, DC 20005-1209			ART UNIT	PAPER NUMBER
			1626	
			NOTIFICATION DATE	DELIVERY MODE
			08/02/2010	ELECTRONIC

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Notice of the Office communication was sent electronically on above-indicated "Notification Date" to the following e-mail address(es):

DBRIPDocket@dbr.com penelope.mongelluzzo@dbr.com

		Application No.	Applicant(s)			
Office Action Summary		10/593,034	SHIMAMOTO ET AL.			
		Examiner	Art Unit			
		MICHAEL BARKER	1626			
Period fo	The MAILING DATE of this communication app or Reply	ears on the cover sheet with the c	orrespondence address			
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
Status						
1) 又	Responsive to communication(s) filed on 24 Ma	av 2010.				
· · · · · · · · · · · · · · · · · · ·	This action is FINAL . 2b) ☐ This action is non-final.					
<i>,</i> —	· · · · · · · · · · · · · · · · · · ·					
- /	closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.					
Dispositi	on of Claims					
)⊠ Claim(s) <u>1-14</u> is/are pending in the application.					
•	4a) Of the above claim(s) is/are withdrawn from consideration.					
· · · · · · · · · · · · · · · · · · ·) Claim(s) is/are allowed.					
· · _ ·	Claim(s) <u>1-3 and 7-9</u> is/are rejected.					
· · · · · · · · · · · · · · · · · · ·	Claim(s) <u>4-6 and 10-14</u> is/are objected to.					
8)[Claim(s) are subject to restriction and/or	election requirement.				
Applicati	on Papers					
9)	The specification is objected to by the Examine	r.				
10)	The drawing(s) filed on is/are: a)☐ acce	epted or b) \square objected to by the E	Examiner.			
	Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).					
	Replacement drawing sheet(s) including the correcti	on is required if the drawing(s) is obj	ected to. See 37 CFR 1.121(d).			
11)☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.						
Priority ι	ınder 35 U.S.C. § 119					
	12)⊠ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a)⊠ All b)□ Some * c)□ None of:					
	1. Certified copies of the priority documents have been received.					
	2. Certified copies of the priority documents have been received in Application No					
	3. Copies of the certified copies of the priority documents have been received in this National Stage					
	application from the International Bureau (PCT Rule 17.2(a)).					
* See the attached detailed Office action for a list of the certified copies not received.						
		·				
Attachmen	t(s)					
1) Notice of References Cited (PTO-892) 4) Interview Summary (PTO-413)						
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) Paper No(s)/Mail Date						
3) Information Disclosure Statement(s) (PTO/SB/08) 5) Notice of Informal Patent Application						
Paper No(s)/Mail Date 6) U Other:						

DETAILED ACTION

By Applicant's amendments submitted 05/24/2010, claims 1-14 are pending in this Application, of which, claim 2 is currently amended. Claims 1-14 were previously rejected under 35 USC 103(a) and Obviousness-type Double Patenting.

Response to Remarks

Applicant traverses the rejection of claims 1-14 under 35 USC 103(a) over WO 03/000698. Applicant states that of the eleven compounds cited from the '698 publication, compounds (1), (2), and (4) have substitutions at the ortho- and metapositions of the benzoyl ring. (6.1, paragraph 3). Applicant's claims are drawn to substitutions only at the para-position. Additionally, Applicant points out compounds (1), (2), and (4) are not radiolabeled. Even excluding these compounds, the remaining eight compounds provide adequate basis to ground this rejection.

Applicant argues there is no suggestion to add a radiolabeled group to the paraposition in particular over the ortho- or meta- positions. The '698 publication specifically teaches a radiolabeled group at the para-position of the benzoyl ring. There need be no suggestion to add a radiolabeled group to the para-position *over* another position.

Instead, the '698 publication need only suggest making modifications to its structurally similar compounds to arrive at the claimed compounds, which it does. Even if the '698 publication states the strongest activity results from the meta-position of the benzoyl ring, it still plainly discloses compounds with a radiolabeled group at the para-position of the benzoyl ring, i.e., compounds claimed by Applicant.

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The '698 publication teaches structurally similar compounds and provides a reason or motivation to modify these compounds to arrive at the claimed compounds. The compounds taught fall within the genus claimed by Applicant and would anticipate Applicant's claims but for the fact Applicant's compounds contain radioactive atoms. In other words, Applicant's claims teach known compounds with the addition of a radiolabel. The '698 publication discloses a valid reason or suggestion to modify the compounds provided at p. 13:

Some compounds of formula (1) are useful for radioisotope labeled ligands for identification of transporter proteins. Isotope labeled ligands may be obtained by well known synthetic procedures, using the hydroxybenzoyl intermediate for R group in the formula (1) with the reaction. for example, of labeled methyl locide to yield the desired labeled ligand as shown in Scheme 2. Some of the radio-isotope labeled methyl iodides are commercially available, including, deuterium-labeled methyl iodide, tritium-labeled methyl iodide, Carbon 14-labeld or Carbon 11-labeled methyl iodides.

As an example of a radiolabeled compound of formula (1), the '698 publication

. This ligand

demonstrates a radiolabeled group at the para-position of the benzoyl ring.

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Applicant states that claim 1 does not recite a radio-labeled methoxy-substituent but instead recites, "3) a straight or branched lower aliphatic alkoxy group",

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Scheme 2 of the '698 publication teaches preparing radio-labeled methoxy-substituent. The Office is reminded that claim 1 does not recite a radio-labeled methoxy-substituent. Instead, claim 1 recites, inter alia, the following radio-labeled substituents:

- 1) a straight or branched lower aliphatic alkyl group;
- 2) a hydroxyl group;
- 3) a straight or branched lower aliphatic alkoxy group;
- 4) an amino group;
- 5) a straight or branched lower aliphatic acylamido group;
- 6) a halogen atom; and
- 7) a straight or branched lower aliphatic haloatkyl group.

(Emphasis added).

Applicant appears to be stating that the language "lower aliphatic alkoxy group", as claimed, does not include methoxy. However, at the paragraph bridging pp. 8 and 9, Applicant's Specification indicates lower aliphatic alkoxy group includes methoxy,

Examples of the radioactive atom possibly contained in the group X in the compound of the formula (1) include 1251, 14C, 3H, 1221, 18F, 12C, 13N and 15O. Specific examples of X having these atoms include methyl group, ethyl group, hydroxyl group, methoxy group ethoxy group, amino group, acetamido group, halogen atoms, halomethyl groups, haloethyl groups and so on. (Emphasis added).

Applicant then argues unexpected advantages at p. 4 of the Response. In doing so, Applicant focuses on the advantages of radiolabeling using ¹²⁵I. This narrow focus ignores the fact that claim 1 encompasses more radioactive atoms than ¹²⁵I alone. Nonetheless, the claimed advantages are stated as follows:

- 1) 123 cmits a gamma ray, which enables direct detection with brain imaging;
- ¹²⁵l can be detected noninvasively in SPECT (single photon emission computed tomography); and
- 3) the binding activity of iodide-substitute (IC₅₀ = 4.8 nM) is higher than the methoxy-substituent (IC₅₀ = 12 nM).⁵

The first two advantages listed are simply properties inherent to ¹²⁵I bestowed on any compound utilizing it as a radiolabel, and while they may be considered advantageous properties of ¹²⁵I, they are not unexpected results. The third advantage demonstrates an improvement in the binding activity of iodide at the para-position versus methoxy as taught in the '698 publication at Table 1. It is not clear, however, that this advantage represents unexpected results or that unexpected results alone would be dispositive of obviousness in this Application.

With regard to the limitation, "a tritium-containing ethyl group", in claim 10, the '698 publication does not teach a tritium containing ethyl group. Instead, the '698 publication discloses, *inter alia*, structurally similar compounds having an alkyl substituent at the benzoyl ring, along with a suggestion that certain of the disclosed compounds could be radiolabeled. This disclosure and suggestion do not render Applicant's claim 10 obvious. With regard to claim 10, this rejection is canceled.

At 6.2 of the Response, Applicant traverses the rejection of claim 9, and states "there is no teaching that would have led a skilled artisan to have identified and used

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the claimed compounds to identify or characterize glutamate transporter proteins." As discussed above, the '698 publication demonstrates structurally similar compounds, suggests modifying them by radiolabeling, and demonstrates a specific example modified by radiolabeling at the para-position of the benzoyl ring. Furthermore, as stated at pp. 6 and 7 of the Non-Final Office Action, "the '698 publication teaches that its radiolabeled ligands are useful for identification of transporter proteins which, as demonstrated throughout the publication, include glutamate transporter proteins."

At 6.3 of the Response, Applicant traverses the rejection of claims 5, 6, 13, and 14. These arguments are persuasive, and this rejection is withdrawn over the process claims, 5, 6, 13, and 14.

At 6.4 of the Response, Applicant traverses the rejection of claims 3, 4, 11, and 12 and argues no reasoning was provided to reject these claims. Scheme 2 of the '698

publication recites the following compound: TBSO

definition of Y at pp. 10 and 11 of the Specification states that when Y is an organometallic group, it may include silicon. It is not clear from claim 3 that Y omits the possibility of the tert-butyldimethylsilyl group disclosed at Scheme 2 of the '698 publication.

In summary, the rejection under 35 USC 103(a) is withdrawn as against claims 4-6 and 10-14 Accordingly, claims 1-3 and 7-9 remain rejected under 35 USC 103(a).

At 7 of the Response, Applicant traverses the rejection over claims 1-14 over US 7247652. Applicant's arguments are persuasive, and this rejection is withdrawn.

Objections

Claims 4-6 and 10-14 are objected to as each claim depends from a rejected base claim.

Conclusion

THIS ACTION IS MADE FINAL. Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the mailing date of this final action.

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Any questions about this Office Action may be directed toward Examiner Michael Barker at 571.272.0303. If, however, attempts to reach Mr. Barker are not successful, the Examiner's supervisor, Joseph McKane, may be reached at 571.272.0699.

/MICHAEL BARKER/ Examiner, Art Unit 1626

/Kamal A Saeed/

Primary Examiner, Art Unit 1626